

**IN THE UNITED STATES PATENT AND
TRADEMARK OFFICE**

Docket No.: **D-6362**

Inventors: **CESARE ET AL.**

Express Mail Label No.: **EG-296014313-US**

Assistant Commissioner for Patents
Box Patent Application
Washington, D.C. 20231

**UTILITY PATENT APPLICATION TRANSMITTAL
UNDER 37 CFR 1.53(b)**

Sir:

Enclosed herewith for filing are:

1. The patent application of:

Inventors: **FRANK C. CESARE;
KAREN A. LEARY and
ANTHONY-J. DiMAIO**

Title: **"LOW MOLECULAR WEIGHT POLYMERS
AND THEIR USE AS DISPERSION AIDS"**

Total pages - 17

2. Application

- a. ☒ New Application
b. ☐ Continuation ☐ Divisional ☐ Continuation-in-part
of U. S. Application Serial No. _____, filed on _____.

3. ☐ sheets of drawings.

4. Oath or Declaration (Total pages - 3)

- a. ☒ Newly executed (original)
b. ☐ Copy from prior application (37 CFR 1.63(d))
(for continuation/divisional application)

5. ☐ Incorporation By Reference

The entire disclosure of the prior application, from which a copy of the oath or declaration is supplied under Box 3 above, is considered as being part of the disclosure of the accompanying application and is hereby incorporated by reference therein.

6. ☒ An assignment of the invention to **UNIROYAL CHEMICAL COMPANY, INC.**, and cover sheet. (Form PTO-1595)

7. ☐ Power of Attorney

8. ☐ Information Disclosure Statement/PTO-1449 (NOT ENCLOSED)

9. ☐ Preliminary Amendment

10. (X) Return Receipt Postcard

11. () Other:

12. The fee has been calculated as shown below:

	Claims Filed	Extra Claims	Rate	Fee
Basic Fee				\$790.00
Total Claims	28-20	8	X \$22	\$176.00
Independent Claims	2-3	0	X \$82	\$0
[] Multiple dependent claims presented			+ \$270	\$0
Total Fee				\$966.00

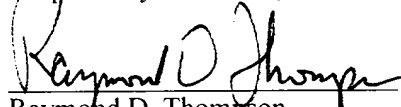
13. (X) Charge Deposit Account No. 21-0525 in the amount of \$ 966.00

14. (X) The Commissioner is hereby authorized to charge payment of the following fees associated with this communication or credit any overpayment to Deposit Account No. 21-0525: any additional filing fees required under 37 CFR 1.16 and any patent application processing fees under 37 CFR 1.17.

15. (X) The Commissioner is hereby authorized to charge payment of the following fees during the pendency of this application or credit any overpayment to Deposit Account No. 21-0525: any patent application processing fees under 37 CFR 1.17 and any filing fees under 37 CFR 1.16 for presentation of extra claims.

Copies in duplicate are enclosed.

Respectfully submitted,



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BY Norma M. Corrigan

DATE August 24, 1998

TITLE OF THE INVENTION
LOW MOLECULAR WEIGHT POLYMERS AND THEIR USE AS DISPERSION
AIDS

5

CROSS-REFERENCES TO RELATED APPLICATIONS
Not applicable.

10

STATEMENT REGARDING FEDERALLY SPONSORED
RESEARCH OR DEVELOPMENT
Not applicable.

BACKGROUND OF THE INVENTION

1. Field of the Invention

15

The present invention relates to low molecular weight polymers formed from monomers comprising ethylene, an alpha-olefin, and optionally a non-conjugated diene, and the use of such polymers to improve the dispersion of reinforcing agents into high molecular weight polymers.

20

2. Description of the Related Art

25

U.S. Patent Nos. 5,391,623 and 5,480,941 are directed to the preparation of masterbatch compositions of elastomers with a high concentration of aramid fibers distributed throughout the elastomer.

U.S. Patent No. 5,527,951 is directed to catalysts for the polymerization of ethylene or the copolymerization of ethylene with alphaolefins and (optionally) nonconjugated polyenes.

30

U.S. Patent No. 5,786,504 is directed to certain catalyst promoters in ethylene polymerization processes.

BRIEF SUMMARY OF THE INVENTION

In one aspect, the present invention relates to a polymer formed from monomers comprising ethylene; $\text{CH}_2=\text{CHQ}$ wherein Q is $\text{C}_1\text{-C}_8$ alkyl; and optionally a non-conjugated polyene; wherein

a) ethylene is present in an amount of from about 67% to about 75% by weight;

b) the non-conjugated polyene is present in an amount of from about 0% to about 30% by weight; and

c) $\text{CH}_2=\text{CHQ}$ is present in an amount of from about 15% to about 40% by weight;
said polymer having a viscosity average molecular weight of from about 4,000 to about 30,000.

In another aspect, the present invention relates to a composition which comprises the polymer described above, and a reinforcing agent.

In yet another aspect, the present invention relates to a composition which comprises:

a) a polymer formed from monomers comprising ethylene; $\text{CH}_2=\text{CHQ}$ wherein Q is $\text{C}_1\text{-C}_8$ alkyl; and optionally a non-conjugated polyene; wherein

i) ethylene is present in an amount of from about 67% to about 75% by weight;

ii) the polyene is present in an amount of from about 0% to about 30% by weight; and

iii) $\text{CH}_2=\text{CHQ}$ is present in an amount of from about 15% to about 40% by weight;

said polymer having a viscosity average molecular weight of from about 4,000 to about 30,000;

b) a reinforcing agent; and

c) a high molecular weight polymer.

In yet another aspect, the present invention relates to a moulded article made from the three part composition described immediately above.

DETAILED DESCRIPTION OF THE INVENTION

The novel low molecular weight polymers of the present invention comprise ethylene, an α -olefin, and optionally a nonconjugated polyene. Preferred

5 α -olefins include propylene, butene-1, pentene-1, hexene-1, 3-methylpentene-1, heptene-1 and octene-1, with propylene being most preferred. Preferred nonconjugated polyenes include dienes such as 5-ethylidene-2-norbornene, 1,4-hexadiene and dicyclopentadiene.

10 Preferably, the polyene component is present in an amount of from about 1% to about 20% by weight, more preferably from about 3% to about 15% by weight. Preferably, the α -olefin component is present in an amount of from about 20% to about 35% by weight, more preferably from

15 about 22% to about 30% by weight. The polymer has a molecular weight (viscosity average) in the range of from about 4,000 to about 30,000, preferably from about 5,000 to about 10,000. Most preferably, the polymer is a solid at room temperature, and yields about 10 mm or less in a

20 needle penetration test as described in Example 1.

The polymerization of the monomers described above may take place in the presence of a catalyst composition which comprises a) a vanadium compound; b) an organo-aluminum compound; and c) a catalyst promoter. Suitable

25 vanadium compounds include vanadium oxytrichloride, vanadium tetrachloride, vanadium acetyl acetonate, vanadyl bis-diethylphosphate, chloro neopentyl vanadate, and the vanadium-containing catalysts described in U.S. Patent No. 5,527,951, the contents of which are

30 incorporated herein by reference.

The organo-aluminum co-catalyst preferably is an alkyl aluminum or an alkyl aluminum halide, with chlorides being the preferred halides. Preferred alkyl aluminum halides include ethyl aluminum sesquichloride,

35 ethyl aluminum dichloride, diethyl aluminum chloride, and

diisobutyl aluminum chloride. Ethyl aluminum sesquichloride and diethyl aluminum chloride are most preferred.

Suitable catalyst promoters include halogenated esters such as butylperchlorocrotonate (BPCC), 4,4,4-trichlorobut-2-eneoate, 2-methyl-4,4,4-trichlorobut-2-eneoate, and other compounds known in the art and described in, for example, U.S. Patent Nos. 5,527,951 and 5,786,504, the contents of which are incorporated herein by reference, with 2-methyl-4,4,4-trichlorobut-2-eneoate (MBEY) being most preferred.

Other suitable supported and unsupported polymerization catalysts would be readily apparent to one of ordinary skill, and include metallocenes, catalytically active titanium, zirconium, hafnium, chromium, etc.

The monomers may be polymerized in the following manner. The catalyst, co-catalyst, promoter, reaction medium and co-monomers are introduced into a reaction vessel. The molar ratio of the catalyst promoter to the vanadium in the vanadium-containing compound is, preferably, in the range of between about 3:1 and about 80:1, more preferably between about 6:1 and about 64:1, and most preferably between about 12:1 and about 48:1.

The molar ratio of the cocatalyst to catalyst plus catalyst promoter is, preferably, in the range of between about 0.5:1 and about 500:1, more preferably between about 1.5:1 and about 100:1, and most preferably between about 2.5:1 and about 10:1. The catalyst concentration can typically range between about 1×10^{-8} and 3×10^{-1} mole of vanadium per liter of total reaction medium.

The reaction medium is an inert medium such as, e.g., pentane, hexane, heptane, octane, isooctane, decane, benzene, toluene and the like, optionally in combination with liquid α -olefins. The polymerization

reaction is typically conducted in the liquid state at a temperature in the range of between about -25°C and about 70°C, for a time which can vary from several minutes to several hours or more depending on the specific reaction conditions and materials, typically between about 15 minutes and 3 hours.

The best reinforcement of elastomeric materials occurs when there is a uniform, non-clumped dispersion of the reinforcing agent(s) in the elastomeric material, and the low molecular weight polymers of the present invention may be used to improve the dispersion of various reinforcing fibers into such high molecular weight polymers. According to the present invention, the reinforcing fibers are dispersed with the low molecular weight polymer to form a reinforcing composition. The low melting characteristic of the preferred low molecular weight polymers allows the liquification and wetting of the reinforcing fibers with much facility. The dispersion may be accomplished by means standard in the art, such as by blending on a rubber mill. The amount of reinforcing material which may be dispersed will vary according to the desired application and the nature of the materials used. In general, it is contemplated that blends of up to about 70% by weight of reinforcing agent will be particularly useful, with about 50% being particularly preferred. The reinforcing composition may subsequently be incorporated into a high molecular weight polymer.

The high molecular weight polymers which may be reinforced according to the present invention include both natural rubber and synthetic rubber compounds. Synthetic rubber compounds include, for example, ethylene/alphaolefin/nonconjugated polyene (EPDM) rubbers, styrene/butadiene rubbers, acrylonitrile/

butadiene (NBR) rubbers, polychloroprene and sulfur modified polychloroprene, polybutadiene rubbers, etc.

Suitable reinforcing agents according to the present invention include aramid fibers (various lengths, short
5 fibers or pulp; for example as disclosed in U.S. Patent No. 5,391,623, the contents of which are incorporated herein by reference), cotton, polyesters, fiberglass, etc.

The masterbatch reinforced high molecular weight
10 polymers of the present invention may be processed by well known means into, e.g., various types of reinforced belts, such as v-belts, timing belts, conveyor belts and drive belts; hoses; seals; diaphragms; cables; roll covers; etc., and may contain other conventional
15 additives such as processing aids, antioxidants, antiozonants, etc.

The following non-limiting examples are illustrative of the processes and products of the present invention.

20

EXAMPLE 1

Preparation of low molecular weight polymer

Into a 3 liter stainless steel stirred autoclave (Buchi, Model BEP 280) with jacketed cooling, a dip tube for feeding ethylene, a thermocouple well, pressure gauge
25 and ports for the introduction of hydrogen, propylene, and the catalyst components, were charged 110 grams of liquid propylene and 8.9 g of 5-ethylidene-2-norbornene (ENB). The temperature was set at 60°C by cooling the jacket with water from a circulating water bath. 25 g of
30 ethylene were then added to the reactor. A solution of 11.1 mmole of ethyl aluminum sesquichloride in 10 ml of hexane from a pressurized bomb was then added to the Buchi followed by enough hydrogen gas to raise the pressure to 270 psig. 43 ml of a hexane solution
35 containing 0.281 mm of vanadium oxytrichloride and 1.31

mmole of MBEY promoter were pumped in continuously over the course of 20 minutes at an inlet pressure of 400 psi. The ensuing exotherm was controlled by the jacket cooling to maintain the temperature of 60°C. The pressure was maintained at 270 psig by feeding ethylene into the Buchi at a rate of 2.4 standard liters per minute to replace the ethylene which was being polymerized. A total of 57.3 g of ethylene was fed in 20 minutes.

The contents of the Buchi were then vented to remove unreacted monomer, and transferred to a two liter agitated pressure vessel (Chemco reactor) containing 200 ml of hexane, 0.1 grams of epoxidized soybean oil and 0.1 g of octadecyl 3,5-di-t-butyl-4-hydroxyhydrocinnamate (Naugard® 76, Uniroyal Chemical Co., Inc., Middlebury, CT) to deactivate the catalyst. The hexane/polymer mixture was then washed twice with 400 ml of deoxygenated water, allowing to settle and decant off the aqueous layer each time. The hexane was then removed by distillation leaving a low molecular weight ethylene-propylene-ENB terpolymer with the following characteristics: $M_v=8,178$; 24.6% propylene; 9.1% ENB; needle penetration (see below): 4.85 mm; yield: 85g; efficiency: 1,759 grams per gram of catalyst.

Three commercial lots of polymer were prepared substantially as described above, and their properties were determined and listed in Table 1 below.

Table 1

Lot	5138/6	5138/7	5138/8
Plant Data			
M_v^1	7190	8210	8180
% ethylene	67	71	71
% diene	11	12.6	13
GPC @ 135°C, ODCB			
Mw	42,000	--	--

	Mn	14,000	--	--
	Mw/Mn	3.0		
	DSC-variable temp.			
	Tg, °C	-46	-41	-40
5	Tm #1, °C	0	21	18
	Tm #2, °C	none	43	43
	DSC @ 190°C			
	OIT, min	4.5		
	Brookfield Vis. (cps, HBT #7)			
10	100°C	74,000	79,000	--
	60°C	670,000	408,000	--

Note: -- means not tested.

15 ¹Mv is a determination of molecular weight by a viscometric method, in which a number average molecular weight is determined by measuring the increase in viscosity of a standard viscosity oil which results when the polymer is dissolved in the oil.

20 The polymers are pale yellow waxy solids at room temperature.

Needle penetration Test

25 Since the hardness of a polymer has some relationship to the crystallinity, molecular weight and tack of the polymer, a needle penetration test can gauge the suitability of the low molecular weight polymers of the present invention. The following is an adaptation of ASTM D 1321-95, "Standard Test Method for Needle
30 Penetration of Petroleum Waxes."

A polymer sample is heated to approximately 100°C in a specimen container (a glass bottle having a one inch inside diameter and a minimum depth of 1.25 inches, filled to a depth of at least 0.75 inches) for one hour
35 or until the sample is homogeneous and free of air

bubbles. The container and contents are cooled to room temperature for 2 hours.

The specimen container is placed on the test shelf of a penetrometer, and the needle is adjusted so that the tip of the needle nearly touches the surface of the specimen. The needle indicator should be in the "zero" position and the total weights of needle and plunger should equal 100 ± 0.15 g. Lock the movable assembly into position.

By means of the fine adjustment knob, the needle tip is brought to just touch the surface of the specimen, watching the reflection of the needle tip as an aid. When in place, the needle shaft is released and held free for 5 ± 0.1 seconds, then re-locked. The indicator shaft is gently depressed until it is stopped by the needle shaft, and the penetration is read from the indicator scale.

The needle is cleaned with hexane to remove any adhering polymer, and the test is repeated three more times, repositioning the needle to a new location each time. The mean of the four penetrations is reported to the nearest 0.1 mm.

EXAMPLE 2

Blending of low molecular weight polymer with aramid fibers

A Brabender internal mixer was warmed to 80°C , and the desired amount of the polymer from Example 1 was added. When the polymer melted, the desired amount of aramid fiber (KEVLAR® merge 1F561 short fiber, DuPont, Wilmington, DE) was added to the melted polymer, the polymer and fibers were mixed for ten minutes at 100 rpm, then allowed to cool and removed. The data for runs A-E are presented below in Table 2. Up to 50% by weight of

aramid fibers were blended with the low molecular weight polymer.

TABLE 2

5		<u>A</u>	<u>B</u>	<u>C</u>	<u>D</u>	<u>E</u>
	Low MW	150 g	150 g	150 g	150 g	140 g
	polymer					
	aramid	45 g	60 g	75 g	112.5 g	140 g
10	fiber					
	% fiber	23%	28.6%	33%	43%	50%

EXAMPLE 3

Triblending of low molecular weight polymer, aramid fiber and high molecular weight EPDM

In this example, four different blends of a low molecular weight polymer according to the present invention, aramid fiber, and a high molecular weight EPDM rubber were made. The low molecular weight polymer was added onto a cold 8" rubber mill. Next, the aramid fiber was added slowly, while increasing the temperature to 150-200° F. The high molecular weight EPDM rubber (Royalene® 521, Uniroyal Chemical Company, Middlebury, CT) was added, the mill was cooled down, and the resulting triblend was stripped off. The various compositions are described in Table 3 below.

TABLE 3

30		<u>Low mw polymer</u>	<u>Aramid fiber</u>	<u>EPDM</u>
	Blend 1	40 g	40 g	85 g
	Blend 2	120 g	40 g	90 g
35	Blend 3	80 g	40 g	80 g

Blend 4 80 g

40 g

80 g

EXAMPLE 4

5 In this example, the rate of incorporation into an
EPDM rubber of aramid fibers alone, and aramid fibers
mixed with a low molecular weight polymer, are compared.
A triblend was prepared as described in Example 3, which
10 contained 33% aramid fiber. As a comparison, a 33%
aramid fiber/ high molecular weight EPDM blend was
prepared as in Example 3, except that the low molecular
weight polymer was omitted. It took 35 minutes to
incorporate the aramid fiber into the high molecular
weight EPDM, compared to 16.5 minutes to incorporate the
15 aramid fiber/low molecular weight polymer into the EPDM,
a time saving of 18.5 minutes (53%).

EXAMPLE 5

20 This example demonstrates a procedure for the
dispersion of aramid fiber (100 grams of 1F 538 Kevlar®)
into 100 grams of Neoprene GNA with the aid of a low
molecular weight polymer (100 grams of Trilene® 77) on a
rubber mill according to the present invention.

25

<u>Time</u>	<u>Temperature</u>	<u>Comments</u>
0	76°F	Start adding Neoprene
3'	94°F	Masticate polymer-no peptizer
6'36"	110°F	Start adding Trilene® 77
30 11'		Start adding Kevlar® slowly
13'30"	108°F	
15'30"		Cool, no heat added, continue adding Kevlar®
18'	123°F	Warms up on addition of fiber
35 19'		Finish adding Kevlar®

21'	150°F	Add heat
23'	190°F	Turn off steam, add cold water
25'	140°F	
27'30"	87°F	Finished, sheet off mill.

5

CLAIMS

1. A polymer formed from monomers comprising ethylene; $\text{CH}_2=\text{CHQ}$ wherein Q is $\text{C}_1\text{-C}_8$ alkyl; and optionally a non-conjugated polyene; wherein

5 a) ethylene is present in an amount of from about 67% to about 75% by weight;

b) the non-conjugated polyene is present in an amount of from about 0% to about 30% by weight; and

10 c) $\text{CH}_2=\text{CHQ}$ is present in an amount of from about 15% to about 40% by weight;
said polymer having a viscosity average molecular weight of from about 4,000 to about 30,000.

2. The polymer of claim 1, wherein the non-
15 conjugated polyene is selected from the group consisting of 5-ethylidene-2-norbornene, 1,4-hexadiene and dicyclopentadiene.

20 3. The polymer of claim 2, wherein Q is methyl.

4. The polymer of claim 3, wherein the polyene component is present in an amount of from about 1% to about 20% by weight.

25 5. The polymer of claim 4, wherein the polyene component is present in an amount of from about 3% to about 15% by weight.

30 6. The polymer of claim 1, wherein the $\text{CH}_2=\text{CHQ}$ component is present in an amount of from about 20% to about 35% by weight.

35 7. The polymer of claim 6, wherein the $\text{CH}_2=\text{CHQ}$ component is present in an amount of from about 22% to about 30% by weight.

8. The polymer of claim 1, wherein the polymer has a viscosity average molecular weight of from about 5,000 to about 10,000.

5 9. The polymer of claim 1, wherein the polymer yields about 10 mm or less in a needle penetration test.

10 10. The polymer of claim 1, wherein said polymer further comprises a reinforcing agent.

11. The polymer of claim 10, wherein the reinforcing agent is selected from the group consisting of aramid fibers, cotton, polyesters, fiberglass, and mixtures thereof.

15 12. The polymer of claim 11, wherein the reinforcing agent comprises aramid fibers.

20 13. The polymer of claim 10, wherein the reinforcing agent is present in an amount of up to about 70% by weight.

14. A composition which comprises:

25 a) a polymer formed from monomers comprising ethylene; $\text{CH}_2=\text{CHQ}$ wherein Q is $\text{C}_1\text{-C}_8$ alkyl; and optionally a non-conjugated polyene; wherein

i) ethylene is present in an amount of from about 67% to about 75% by weight;

30 ii) the polyene is present in an amount of from about 0% to about 30% by weight; and

iii) $\text{CH}_2=\text{CHQ}$ is present in an amount of from about 15% to about 40% by weight;

said polymer having a viscosity average molecular weight of from about 4,000 to about 30,000;

35 b) a reinforcing agent; and

c) a high molecular weight polymer.

15. The composition of claim 14, wherein the non-conjugated polyene is selected from the group consisting of 5-ethylidene-2-norbornene, 1,4-hexadiene and dicyclopentadiene.

16. The composition of claim 15, wherein Q is methyl.

17. The composition of claim 16, wherein the polyene component is present in an amount of from about 1% to about 20% by weight.

18. The composition of claim 17, wherein the polyene component is present in an amount of from about 3% to about 15% by weight.

19. The composition of claim 14, wherein the $\text{CH}_2=\text{CHQ}$ component is present in an amount of from about 20% to about 35% by weight.

20. The composition of claim 19, wherein the $\text{CH}_2=\text{CHQ}$ component is present in an amount of from about 22% to about 30% by weight.

21. The composition of claim 14, wherein the polymer of part a) has a viscosity average molecular weight of from about 5,000 to about 10,000.

22. The composition of claim 14, wherein the polymer of part a) yields about 10 mm or less in a needle penetration test.

23. The composition of claim 14, wherein the reinforcing agent is selected from the group consisting of aramid fibers, cotton, polyesters, fiberglass, and mixtures thereof.

5

24. The composition of claim 23, wherein the reinforcing agent comprises aramid fibers.

25. The composition of claim 14, wherein the high molecular weight polymer is selected from the group consisting of natural rubber and synthetic rubber.

26. The composition of claim 25, wherein the synthetic rubber is selected from the group consisting of ethylene/alphaolefin/nonconjugated polyene (EPDM) rubbers, styrene/butadiene rubbers, acrylonitrile/butadiene (NBR) rubbers, polychloroprene and sulfur modified polychloroprene, and polybutadiene rubbers.

27. A moulded article made from the composition of claim 14.

28. The article of claim 27, wherein the article is selected from the group consisting of a v-belt, a timing belt, a conveyor belt, a drive belt, a hose, a seal, a diaphragm, a cable and a roll cover.

ABSTRACT OF THE DISCLOSURE

Disclosed are low molecular weight polymers formed from monomers comprising ethylene, an alpha-olefin, and optionally a non-conjugated diene, and the use of such
5 polymers to improve the dispersion of reinforcing agents into high molecular weight polymers.

**COMBINED DECLARATION FOR PATENT
APPLICATION AND POWER OF ATTORNEY**

**Atty Docket No.
D-6362**

**(Includes Reference to PCT International
Applications)**

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am an original, first and joint inventor of the invention entitled "**LOW MOLECULAR WEIGHT POLYMERS AND THEIR USE AS DISPERSION AIDS**" which is described and claimed in the patent specification which

(X) is attached hereto,

() was filed on and accorded serial number ,
and for which invention Letters Patent are sought.

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, Sec. 1.56(a).

I hereby claim foreign priority benefits under Title 35, United States Code, Sec. 119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed:

PRIOR FOREIGN/PCT APPLICATIONS AND PRIORITY CLAIMS UNDER 35 U.S.C. 119			
COUNTRY (if PCT, indicate "PCT")	APPLICATION NO.	DATE OF FILING (month, day, year)	PRIORITY CLAIMED

**COMBINED DECLARATION FOR PATENT
APPLICATION AND POWER OF ATTORNEY**

**Atty Docket No.
D-6362**

**(Includes Reference to PCT International
Applications)**

Page 2

I hereby claim the benefit under Title 35, United States Code, Sec.120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, Sec.112, I acknowledge the duty to disclose material information as defined in Title 37, Code Federal Regulation Sec.1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

PRIOR U.S. APPLICATIONS OR PCT INTERNATIONAL APPLICATIONS DESIGNATING THE U.S. FOR BENEFIT UNDER 35 U.S.C. 120					
U.S. APPLICATIONS			STATUS (CHECK ONE)		
US APPLN. NO.	US FILING DATE		PATENTED	PENDING	ABANDONED
PCT APPLICATIONS DESIGNATING THE U.S.					
PCT APPLICATION NO.	PCT FILING DATE	US SERIAL NOS. ASSIGNED (if any)			

I hereby appoint the following attorney's and/or agent(s) to prosecute this application and to transact all business in the Patent and Trademark Office connected therewith: **Raymond D. Thompson - Reg. No. 30,695, and Daniel Reitenbach - Reg. No. 30,970.**

Address all correspondence to: Raymond D. Thompson Uniroyal Chemical Company, Inc. World Headquarters Middlebury, CT 06749	Direct telephone calls to: Raymond D. Thompson tel. no.: (203) 573-4385.
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COMBINED DECLARATION FOR PATENT
APPLICATION AND POWER OF ATTORNEY

Atty Docket No.
D-6362

(Includes Reference to PCT International Applications)

Page 3

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

2 0 1	FULL NAME OF INVENTOR	FRANK C. CESARE		
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	SIGNATURE	<i>Frank C. Cesare</i>	DATE	<i>Aug 24, 1998</i>

2 0 2	FULL NAME OF INVENTOR	KAREN A. LEARY		
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	SIGNATURE	<i>Karen A. Leary</i>	DATE	<i>August 24, 1998</i>

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